

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C08F 210/18, 2/06, 4/645	A1	(11) International Publication Number: WO 99/41294 (43) International Publication Date: 19 August 1999 (19.08.99)
(21) International Application Number: PCT/US99/03264		(81) Designated States: BR, CA, CN, JP, KR, MX, SG, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).
(22) International Filing Date: 17 February 1999 (17.02.99)		
(30) Priority Data: 60/074,814 17 February 1998 (17.02.98) US		Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(71) Applicant: EXXON CHEMICAL PATENTS INC. [US/US]; 5200 Bayway Drive, Baytown, TX 77520-5200 (US).		
(72) Inventors: SCHIFFINO, Rinaldo, S.; 4809 Threadneedle, Wilmington, DE 19807 (US). CROWTHER, Donna, J.; 1104 Cactus Drive, Baytown, TX 77521 (US).		
(74) Agents: MULLER, William, G. et al.; Exxon Chemical Company, P.O. Box 2149, Baytown, TX 77522-2149 (US).		

(54) Title: ETHYLENE COPOLYMERIZATION PROCESS

(57) Abstract

This invention is a solution process for the preparation of ethylene- α -olefin-diolefin copolymers comprising contacting ethylene, one or more α -olefin monomer, and one or more cyclic diene monomer, with a catalyst composition comprising a bridged, bis(cyclopentadienyl) zirconium compound having an unsubstituted cyclopentadienyl ligand, a multiply substituted cyclopentadienyl ligand, said ligands bridged by a covalent bridging group containing one or more Group 14 element, and two uninegative, activation reactive ligands and a catalyst activator compound. The invention process exhibits high catalyst activity, high comonomer incorporation and high diene monomer conversion rates and is particularly suitable for the preparation of elastomeric ethylene-propylene or ethylene-propylene-diene monomer elastomers.

BEST AVAILABLE COPY

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

TITLE

Ethylene Copolymerization Process

5

TECHNICAL FIELD

This invention relates to the preparation of ethylene- α -olefin polymers under solution polymerization conditions using catalyst compositions based on biscyclopentadienyl derivatives of zirconium.

10

BACKGROUND OF THE INVENTION

Polymers comprising ethylene and at least one or more α -olefin and optionally one or more diolefin make up a large segment of polyolefin polymers and will be addressed for convenience as "ethylene- α -olefin-diolefin copolymers" herein. Such polymers range from crystalline polyethylene copolymers to largely amorphous elastomers, with a new area of semi-crystalline "plastomers" in between. In particular, ethylene- α -olefin-diolefin elastomers are a well established class of industrial polymers having a variety of uses associated with their elastomeric properties, their thermo-oxidative stability, their solubility in hydrocarbon oleaginous fluids, and their capability for modifying the properties of polyolefin blends. Included in this terminology are the commercially available EPM (ethylene-propylene monomer) and EPDM (ethylene-propylene-diene monomer) rubbery polymers, both being vulcanizable by cross-linking, the addition of the diolefin, also known as diene monomer, providing increased ease of both cross-linking and functionalization.

25

Commercially prepared ethylene- α -olefin-diolefin elastomers have been traditionally been made via Ziegler-Natta polymerization with homogenous catalyst compositions largely based on vanadium or titanium. Newer metallocene catalyst compounds have received attention due to their ease of larger monomer incorporation and potential increases in polymerization activities. U.S. patent 5,324,800 describes

30

compounds have received attention due to their ease of larger monomer incorporation and potential increases in polymerization activities. U.S. patent 5,324,800 describes metallocenes having substituted and unsubstituted cyclopentadienyl ligands which are suitable for producing high molecular weight olefin polymers, including linear, low density copolymers of ethylene with minor amounts of α -olefin. WO95/277147 describes bridged and unbridged Group 4 metallocene compounds wherein the cyclopentadienyl ligands have two or four adjacent substituents forming one or two alkylenic cycles of from 4 to 8 carbon atoms. These compounds are said to be useful for ethylene copolymerization and propylene polymerization, including elastomeric copolymers of ethylene, α -olefins and non-conjugated diolefins. Ethylene copolymerization with propylene is reported in examples 28-30 and in Table 3. U.S. patent 5,543,373 describes bridged metallocenes having two differing π ligands which are said to be of high activity. Copolymers of ethylene with 1-olefins and/or one or more diene monomers are produced in a preferred process according to the invention. Example R illustrates an ethylene-propylene-diene terpolymer rubber prepared with dimethylsilanediyl(2-methyl-4-phenyl-1-indenyl)(2,3,5-trimethyl-1-cyclopentadienyl) zirconium dichloride.

Ethylene copolymerization is described in WO 95/27717 with zirconocenes having a Cp cyclopentadienyl group with one or two alkylenic cycles of 4 to 8 carbon atoms and a Cp' cyclopentadienyl group having up to 4 R substituents. Example 12 illustrates the preparation of isopropyliden(cyclopentadienyl) (2,3-cyclotetramethyleneinden-1-yl) zirconium dichloride. Example 19 illustrates syndiotactic propylene polymerization with this catalyst.

25

A high temperature solution process for the preparation of ethylene- α -olefin copolymers is described in EP-A-0 612 769. The catalyst compositions are based on bis(cyclopentadienyl/indenyl/fluorenyl) titanocenes/zirconocenes which are combined with an alkyl aluminum compound and an ionizing ionic compound which provides a non-coordinating anion. Asymmetrically substituted catalysts are illustrated. The

compound is reacted with the organoaluminum compound, then reacted with ionizing ionic compound, and subsequently added to the polymerization reactor. High molecular weight polymer is said to be produced at high efficiency.

5 A high activity supported catalyst suitable for ethylene copolymers is described in U.S. patent 5,240,894. The catalyst isopropylidene(cyclopentadienyl)(fluorenyl) zirconium dichloride is a preferred metallocene embodiment. Example 10 illustrates an ethylene-propylene copolymerization.

10 It remains important in industry to develop efficient copolymerization processes, and in particular, those capable of high productivity of polymer per unit weight of catalyst compound.

INVENTION DISCLOSURE

15 The invention is a polymerization process for the preparation of ethylene- α -olefin-diolefin copolymers comprising contacting ethylene, one or more α -olefin monomer, and optionally, one or more cyclic diolefin monomer, with a catalyst composition prepared from at least one a catalyst activator and at least one bridged, bis(cyclopentadienyl) zirconium compound having an unsubstituted cyclopentadienyl
20 ligand, a bulky, substituted cyclopentadienyl ligand, said ligands bridged by a covalent bridging group containing one or more Group 14 element, said process conducted in a solution polymerization process. The invention process exhibits high catalyst activity, high comonomer incorporation and high diene monomer conversion rates.

25 BEST MODE AND EXAMPLES OF THE INVENTION

30 The ethylene- α -olefin-diolefin copolymers of this invention (hereinafter referred to as "EPC") is meant to include copolymers, terpolymers, tetrapolymers, etc. It typically comprises ethylene, one or more alpha-olefins, and optionally, one or more cyclic diolefin monomers; it is typically substantially amorphous; and it will typically have a substantially random arrangement of at least the ethylene and the alpha-olefin

cyclic diolefin monomers; it is typically substantially amorphous; and it will typically have a substantially random arrangement of at least the ethylene and the alpha-olefin monomers. Thus both of ethylene-containing elastomer and plastomer copolymers can be prepared by the invention process.

5

The EPC capable of preparation in accordance with the invention process generally can have a molecular weight range typically between about 20,000 and up to about 500,000 or higher, more typically between about 60,000 and 300,000 where the molecular weight is number-average ("M_n").

10

Typically elastomeric EPC is "substantially amorphous", and when that term is used to define the EPC elastomers of this invention it is to be taken to mean having a degree of crystallinity less than about 25% as measured by means known in the art, preferably less than about 15%, and more preferably less than about 10%. The three major known methods of determining crystallinity are based on specific volume, x-ray diffraction, and infrared spectroscopy. Another well-established method, based on measurement of heat content as a function of temperature through the fusion range, is carried out using differential scanning calorimetric measurements. It is known that these independent techniques lead to reasonably good experimental agreement. The degree of randomness of the arrangement of monomers in the EPC elastomeric polymers also affects the crystallinity and is appropriately characterized by the degree of crystallinity.

15

20

25

30

Additionally, it is known in the art that the tendency of a particular combination of catalyst composition and monomers to produce blocky, random, or alternating polymers can be characterized by the product of the reactivity ratios defined for the given monomers under the specific reaction conditions encountered. If this product is equal to 1.0, the sequence distribution will be perfectly random; the more the product is less than 1.0, the more the monomers will tend to have a "blocky" sequence distribution. Generally speaking, the segments of a polymer which crystallize are

linear segments of a polymer which have a number of identical (both by chemical make-up and stereo-specific orientation) units in a row. Such segments are said to be "blocky". If there is little or no such sequential order within the segments making up a polymer chain, that chain will be very unlikely to conform itself into the correct shape to fit into the spatial order of a crystal and will accordingly exhibit a low degree of crystallinity. See, "Ethylene-Propylene Copolymers. Reactivity Ratios, Evaluation and Significance", C. Cozewith and G. Ver Strate, *Macromolecules*, Vol. 4, No. 4, 482-489 (1971). The EPC elastomers of this invention accordingly can be characterized in one embodiment by the limitation that its method for preparation has a reactivity ratio product less than 2.0, preferably less than about 1.5, and more preferably less than about 1.25.

The EPC elastomers of the invention will contain about 10 to about 91 weight percent ethylene, preferably about 20 to 88 weight percent ethylene. The EPC elastomers of the invention preferably contain from 35 to 75 weight percent ethylene.

The α -olefins suitable for use in the preparation of the EPC, or for the polyethylene copolymers, are preferably C_3 to C_{20} α -olefins, but will include higher carbon number olefins such as polymerizable macromers having up to five hundred carbon atoms, or more. Illustrative non-limiting examples of such α -olefins are one or more of propylene, 1-butene, 1-pentene, 1-hexene, 1-octene, and 1-decene. Included in the term α -olefins for the purposes of describing effectively copolymerized monomers are the constrained-ring cyclic monoolefins such as cyclobutene, cyclopentene, norbornene, alkyl-substituted norbornenes, alkenyl-substituted norbornenes, and the higher carbon number cyclic olefins known in the art, see U.S. patent 5,635,573, incorporated herein by reference for purposes of U.S. patent practice. The α -olefin content of the EPC ranges depending upon selection of the specific α -olefin or α -olefins, being more for lower carbon number monomers, for example, about 10 to about 91 wt.%, preferably about 20 to about 88 wt.% for propylene; and, 5 to 35 mol.%, preferably 7.5 to 25 mol.% and most preferably 10 to 20 mol.% for 1-

octene. The EPC elastomers typically have above about 25 mol.% α -olefin incorporation. For the more crystalline polyethylene copolymers the range of comonomer incorporation will typically be below 25 mol.% and more typically below about 15 mol.%. In terms of polymer density, the elastomers are typically below about 0.860 g/cm³, and the plastomer ethylene copolymers are from about 0.860 to 0.915.

10 The diene monomers, or diolefins, useful in this invention include those typically used in known EPDM polymers. The typically used diene monomers are

generally selected from the cyclic diolefins having about 6 to about 15 carbon atoms, for example:

- A. single ring alicyclic dienes such as 1,3-cyclopentadiene; 1,4-cyclohexadiene; 1,5-cyclooctadiene and 1,5-cyclododecadiene;
- B. multi-ring alicyclic fused and bridged ring dienes such as tetrahydroindene, methyl tetrahydroindene, dicyclopentadiene; bicyclo-(2,2,1)-hepta-2,5-diene; alkenyl, alklinidene, cycloalkenyl and cycloalkylidene norbornenes such as 5-methylene-2-norbornene (MNB), 5-ethylidene-2-norbornene, 5-isopropylidene-2-norbornene, 5-(4-cyclopentenyl)-2-norbornene, 5-cyclohexylidene-2-norbornene, 5-vinyl-2-norbornene; and
- C. cycloalkenyl-substituted alkenes, such as allyl cyclohexene, vinyl cyclooctene, allyl cyclohexene, vinyl cyclooctene, allyl cyclodecene, vinyl cyclododecene.

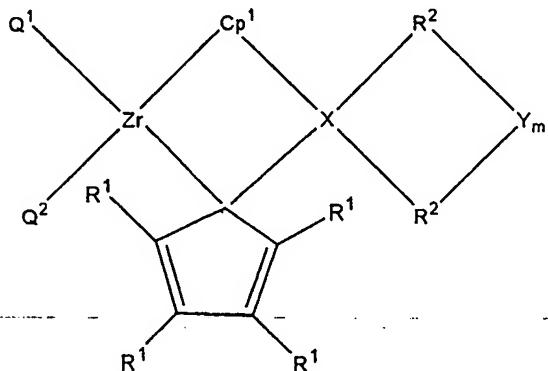
25 Of these, the preferred dienes are dicyclopentadiene, 5-methylene-2-norbornene, 5-ethylidene-2-norbornene and 5-vinyl-2-norbornene. It will be apparent that a mix of such dienes can also be utilized. The content of the diene monomer in the EPC elastomer can be 0 to about 20 weight percent, and if used, preferably 0.5 to about 30 15 weight percent, and most preferably about 2.0 to about 12.0 weight percent.

Surprisingly, diene incorporation greater than 5.0 wt.%, even greater than 8.0 wt.% is made possible using the process of this invention.

The bridged, bis(cyclopentadienyl) compounds of this invention typically comprise those having ancillary ligands including the unsubstituted cyclopentadienyl ligand, a multiply substituted, bulky cyclopentadienyl ligand, the cyclopentadienyl ligands covalently bridged, and two uninegative, activation reactive ligands at least one of which that can be abstracted for activation of the remaining metal compound to a catalytically active state and one of which is either similarly abstractable or has a σ -bond to the transition metal into which an olefin or diolefin can insert for coordination polymerization.

Thus the bridged, bis(cyclopentadienyl) zirconium compounds of the present invention have the following structure:

15



20

where the unsubstituted cyclopentadienyl ring is Cp^1 , and the substituted cyclopentadienyl ring (Cp^2R_n) has at least two non-hydrogen radical substituents R^1 , wherein each R^1 is, independently, a radical selected from:

- hydrogen radical,
- hydrocarbyl, silyl or germyl radicals having from 1 to 20 carbon, silicon or germanium atoms,

c) substituted hydrocarbyl, silyl or germyl radicals as defined, wherein one or more hydrogen atoms is replaced by a halogen radical, an amido radical, a phosphido radical, an alkoxy radical, an aryloxy radical or any other radical containing a Lewis acidic or basic functionality,

5 d) C₁-C₂₀ hydrocarbyl-substituted metalloid radicals wherein the metalloid is selected from the group consisting of Ge, Sn and Pb,

e) halogen radicals,

f) amido radicals,

g) phosphido radicals,

10 h) alkoxy radicals, and

i) alkylborido radicals; or

at least two R¹ groups are joined together to form, along with the carbon atoms to which they are attached, a C₄-C₂₀ ring structure, which is saturated or partially saturated, and substituted or unsubstituted, the ring structure substitution being selected 15 from one or more R¹ groups as defined under (a) – (i) above;

X is selected from the group consisting of C, Si, Ge, Sn and Pb;

R² is selected from the group consisting of H, unsubstituted C₁-C₂₀ hydrocarbyl, C₁-C₂₀ hydrocarbyl substituted with at least one Si, and C₁-C₂₀ hydrocarbyl substituted with at 20 – least one Ge;

m = 0 or 1,

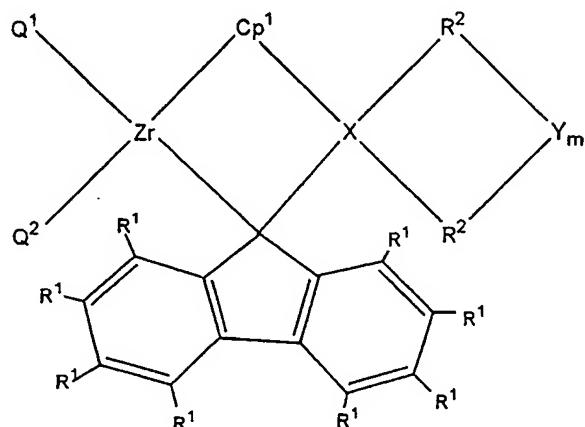
25 Y is selected from the group consisting of hydrocarbyl radicals, hydrosilyl radicals, and hydrogermyl radicals; and

Q¹ and Q² are non-cyclopentadienyl radicals wherein

30 i) independently, each Q is selected from the group consisting of halide, hydride, unsubstituted C₁-C₂₀ hydrocarbyl, C₁-C₂₀ hydrocarbyl substituted

ii) together, Q¹ and Q² may form an alkylidene, cyclometallated hydrocarbyl or any other divalent anionic chelating ligand.

Preferred zirconium compounds where two sets of R¹ groups are joined together has the following structure:



where the unsubstituted cyclopentadienyl ring is Cp¹, and the substituted fluorenyl ring has at least two non-hydrogen radical substituents, wherein each R¹ is, independently, a radical selected from:

- a) hydrogen radical,
- b) hydrocarbyl, silyl or germyl radicals having from 1 to 20 carbon, silicon or germanium atoms,
- c) substituted hydrocarbyl, silyl or germyl radicals as defined, wherein one or more hydrogen atoms is replaced by a halogen radical, an amido radical, a phosphido radical, an alkoxy radical, an aryloxy radical or any other radical containing a Lewis acidic or basic functionality,
- d) C₁-C₂₀ hydrocarbyl-substituted metalloid radicals wherein the metalloid is selected from the group consisting of Ge, Sn and Pb,
- e) halogen radicals,
- f) amido radicals,

- e) halogen radicals,
- f) amido radicals,
- g) phosphido radicals,
- h) alkoxy radicals, and
- 5 i) alkylborido radicals; or

at least two R¹ groups are joined together to form, along with the carbon atoms to which they are attached, a C₄-C₂₀ ring structure, which is saturated or partially saturated, and substituted or unsubstituted, the ring structure substitution being selected from one or more R¹ groups as defined under (a) – (i) above;

10

X is selected from the group consisting of C, Si, Ge, Sn and Pb;

15 R² is selected from the group consisting of H, C₁-C₂₀ hydrocarbyl, C₁-C₂₀ hydrocarbyl

substituted with at least one Si, and C₁-C₂₀ hydrocarbyl substituted with at least one

Ge;

m = 0 or 1,

20 Y is selected from the group consisting of hydrocarbyl radicals, hydrosilyl radicals, and

hydrogermyl radicals; and

Q¹ and Q² are non-cyclopentadienyl radicals wherein

- i) independently, each Q is selected from the group consisting of halide, hydride, unsubstituted C₁-C₂₀ hydrocarbyl, C₁-C₂₀ hydrocarbyl substituted with at 25 least one R¹ group as defined under (a) – (i) above, alkoxide, aryloxide, amide, halide or phosphide; or
- ii) together, Q¹ and Q² may form an alkylidene, cyclometallated hydrocarbyl or any other divalent anionic chelating ligand.

Such compounds can also include an L_w complexed thereto wherein L is a neutral Lewis base such as diethylether, tetrahydrofuran, dimethylaniline, aniline, trimethylphosphine, n-butylamine, and the like; and "w" is a number from 0 to 3.

5 The term "cyclopentadienyl" refers to a 5-member ring having delocalized bonding within the ring and typically being bound to M through η^5 -bonds, carbon typically making up the majority of the 5-member positions.

10 Examples of the bridged, bis(cyclopentadienyl) zirconium compounds of the invention include:

dimethylsilanyl(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dichloride,

15 dimethylsilanyl(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dimethyl,

dimethylsilanyl(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dibenzyl,

20 diphenylsilanyl(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dichloride,

diphenylsilanyl(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dimethyl,

diphenylsilanyl(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dibenzyl,

25 diphenylmethylene(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dichloride,

methylphenylmethylene(cyclopentadienyl)(trimethylcyclopentadienyl)
zirconium dimethyl,

diphenylmethylene(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
30 dibenzyl,

silacyclobutyl(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dichloride,
silacyclobutyl(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dimethyl,
5 silacyclobutyl(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dibenzyl,
isopropylidene(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dichloride,
isopropylidene(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
10 dimethyl,
isopropylidene(cyclopentadienyl)(trimethylcyclopentadienyl) zirconium
dibenzyl,
dimethylsilanyl(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium
dichloride,
15 dimethylsilanyl(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium
dimethyl,
dimethylsilanyl(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium
dibenzyl,
diphenylsilanyl(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium
20 dichloride,
diphenylsilanyl(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium
dimethyl,
diphenylsilanyl(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium
dibenzyl,
25 diphenylmethylene(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium
dichloride,
methylphenylmethylene(cyclopentadienyl)(tetramethylcyclopentadienyl)
zirconium dimethyl,
diphenylmethylene(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium
30 dibenzyl,

silacyclobutyl(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium

dichloride,

silacyclobutyl(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium

dimethyl,

5 silacyclobutyl(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium

dibenzyl,

isopropylidene(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium

dichloride,

isopropylidene(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium

10 dimethyl,

isopropylidene(cyclopentadienyl)(tetramethylcyclopentadienyl) zirconium

dibenzyl.

15 Substituted versions where a hydride, hydrocarbyl, germyl or silyl group replaces one or both chloride ligands of M are suitable in accordance with invention particularly where ionizing anion precursors are activators. Separate or *in situ* alkylation is typical, e.g., dimethyl replacing dichloride.

20 A preferred catalyst according to the invention will be those in which Cp² is tri- or tetra-alkyl substituted with methyl, ethyl, isopropyl or tertiary butyl alkyl groups, or mixed combinations of two or more such alkyl groups. Preferred bridging groups, those particularly suited for increased activity and increased molecular weights, are both silacyclic and aryl-group containing methylene groups, e.g., silacyclobutyl, methylphenylmethylene and diphenylmethylene.

25 The bridged, bis(cyclopentadienyl) zirconium compounds according to the invention may be activated for olefin polymerization catalysis in any manner sufficient both to remove or complex one Q group such that the metal center becomes sufficiently electron deficient to attract olefinically unsaturated monomers and such that the other Q bond is either itself, or is abstracted and substituted with another Q

bond, sufficiently weak so as to permit insertion of it into the olefinically unsaturated monomer to yield a growing polymer, in the manner of traditional coordination/insertion polymerization. The traditional activators of metallocene polymerization art are suitable, those typically include Lewis acids such as aluminum alkyls or alumoxane compounds, and ionizing anion pre-cursors that abstract one Q so as ionize the metal center into a cation and provide a counter-balancing noncoordinating anion.

The term "noncoordinating anion" means an anion which either does not coordinate to said transition metal cation or which is only weakly coordinated to said cation thereby remaining sufficiently labile to be displaced by a neutral Lewis base. "Compatible" noncoordinating anions are those which are not degraded to neutrality when the initially formed complex decomposes. Further, the anion will not transfer an anionic substituent or fragment to the cation so as to cause it to form a neutral four coordinate metallocene compound and a neutral by-product from the anion. Noncoordinating anions useful in accordance with this invention are those which are compatible, stabilize the metallocene cation in the sense of balancing its ionic charge in a +1 state, yet retain sufficient lability to permit displacement by an ethylenically or acetylenically unsaturated monomer during polymerization. Additionally, the anions useful in this invention will be large or bulky in the sense of sufficient molecular size to largely inhibit or prevent neutralization of the metallocene cation by Lewis bases other than the polymerizable monomers that may be present in the polymerization process. Typically the anion will have a molecular size of greater than or equal to about 4 angstroms.

25

Descriptions of ionic catalysts for coordination polymerization comprised of metallocene cations activated by ionizing anion pre-cursors appear in the early work in EP-A-0 277 003, EP-A-0 277 004, US patents 5,198,401 and 5,278,119, and WO92/00333. These teach a preferred method of preparation wherein metallocene compounds are protonated by an anionic precursors such that an alkyl/hydride group is

abstracted from a transition metal to make it both cationic and charge-balanced by the non-coordinating anion. The use of ionizing ionic compounds not containing an active proton but capable of producing the both the active metallocene cation and an noncoordinating anion is also known. See, EP-A-0 426 637, EP-A- 0 573 403 and 5 U.S. patent 5,387,568. Reactive cations other than Bronsted acids capable of ionizing the metallocene compounds include ferrocenium triphenylcarbonium and triethylsilylinium cations. Any metal or metalloid capable of forming a coordination complex which is resistant to degradation by water (or other Bronsted or Lewis Acids) may be used or contained in the anion of the second activator compound. Suitable 10 metals include, but are not limited to, aluminum, gold, platinum and the like. Suitable metalloids include, but are not limited to, boron, phosphorus, silicon and the like. The description of non-coordinating anions and precursors thereto of these documents are incorporated by reference for purposes of U.S. patent practice.

15 An additional method of making the ionic catalysts uses ionizing anion precursors which are initially neutral Lewis acids but form the cation and anion upon ionizing reaction with the metallocene compounds, for example *tris*(pentafluorophenyl) boron acts to abstract an alkyl, hydride or silyl ligand to yield a metallocene cation and stabilizing non-coordinating anion, see EP-A-0 427 697 and EP-A-0 520 732. Ionic 20 catalysts for addition polymerization can also be prepared by oxidation of the metal centers of transition metal compounds by anionic precursors containing metallic oxidizing groups along with the anion groups, see EP-A-0 495 375. The description of non-coordinating anions and precursors thereto of these documents are similarly incorporated by reference for purposes of U.S. patent practice.

25 Examples of suitable anion precursors capable of ionic cationization of the metallocene compounds of the invention, and consequent stabilization with a resulting noncoordinating anion include trialkyl-substituted ammonium salts such as are well-known in the art, see U.S. patent 5,198,401 and WO-A-96\33227, and others above.

Further examples of suitable anion precursors include those comprising a stable carbonium ion, and a compatible non-coordinating anion. These include:

5 tropillium tetrakis(pentafluorophenyl) borate,
triphenylmethylium tetrakis(pentafluorophenyl) borate,
benzene (diazonium) tetrakis(pentafluorophenyl) borate,
tropillium phenyltris(pentafluorophenyl) borate,
triphenylmethylium phenyl-trispentafluorophenyl borate,
benzene (diazonium) phenyltrispentafluorophenyl borate,
10 tropillium tetrakis (2,3,5,6-tetrafluorophenyl) borate,
triphenylmethylium tetrakis (2,3,5,6-tetrafluorophenyl) borate,
benzene (diazonium) tetrakis (3,4,5-trifluorophenyl) borate,
tropillium tetrakis (3,4,5-trifluorophenyl) borate,
benzene (diazonium) tetrakis (3,4,5-trifluorophenyl) borate,
15 tropillium tetrakis (3,4,5-trifluorophenyl) aluminate,
triphenylmethylium tetrakis (3,4,5-trifluorophenyl) aluminate,
benzene (diazonium) tetrakis (3,4,5-trifluorophenyl) aluminate,
tropillium tetrakis (1,2,2-trifluoroethenyl) borate,
triphenylmethylium tetrakis (1,2,2-trifluoroethenyl) borate,
20 benzene (diazonium) tetrakis (1,2,2-trifluoroethenyl) borate, tropillium tetrakis
(2,3,4,5-tetrafluorophenyl) borate,
triphenylmethylium tetrakis (2,3,4,5-tetrafluorophenyl) borate,
benzene (diazonium) tetrakis (2,3,4,5-tetrafluorophenyl) borate,
and, diethylanilinium n-butyltris(pentafluorophenyl) borate.

25

Where the metal ligands include Q halide moieties, such as in (cyclopentadienyl)dimethylsilyl(tetramethylcyclopentadienyl) zirconium dichloride, which are not capable of discrete ionizing abstraction under standard conditions, these moieties can be converted via known alkylation reactions with organometallic compounds such as lithium or aluminum hydrides or alkyls, alkylalumoxanes,

Grignard reagents, etc. See EP-A-0 500 944, EP-A1-0 570 982 and EP-A1-0 612 769 for processes describing the reaction of alkyl aluminum compounds with dihalide substituted metallocene compounds prior to or with the addition of activating anion precursor compounds.

5

Known alkylalumoxanes are additionally suitable as catalyst activators, particularly for the invention metal compounds comprising the halide ligands. The alumoxane component useful as catalyst activator typically is an oligomeric aluminum compound represented by the general formula $(R-Al-O)_n$, which is a cyclic compound, or $R(R-Al-O)_nAlR_2$, which is a linear compound. In the general alumoxane formula R is a C₁ to C₅ alkyl radical, for example, methyl, ethyl, propyl, butyl or pentyl and "n" is an integer from 1 to about 50. Most preferably, R is methyl and "n" is at least 4. Alumoxanes can be prepared by various procedures known in the art. For example, an aluminum alkyl may be treated with water dissolved in an inert organic solvent, or it may be contacted with a hydrated salt, such as hydrated copper sulfate suspended in an inert organic solvent, to yield an alumoxane. Generally, however prepared, the reaction of an aluminum alkyl with a limited amount of water yields a mixture of the linear and cyclic species of the alumoxane.

20

When using ionic catalysts comprising bridged Group 4 metal cations and non-coordinating anions, the total catalyst composition will generally additionally comprise one or more scavenging compounds. The term "scavenging compounds" as used in this application and its claims is meant to include those compounds effective for removing polar impurities from the reaction solvent. Such impurities can be inadvertently introduced with any of the polymerization reaction components, particularly with solvent, monomer and catalyst feed, and adversely affect catalyst activity and stability. It can result in decreasing or even elimination of catalytic activity, particularly when a metallocene cation-noncoordinating anion pair is the catalyst composition. The polar impurities, or catalyst poisons include water, oxygen, metal impurities, etc. Preferably steps are taken before provision of such into the

25

30

reaction vessel, for example by chemical treatment or careful separation techniques after or during the synthesis or preparation of the various components, but some minor amounts of scavenging compound will still normally be required in the polymerization process itself. Typically the scavenging compounds will be an organometallic compound such as the Group-13 organometallic compounds of 5,153,157, 5,241,025, EP-A-0 426 638 and WO-A-91/09882 and WO-A-94/03506, noted above, and those of WO-A-93/14132 and WO-A-97/22635. Exemplary compounds include triethyl aluminum, triethyl borane, tri-isobutyl aluminum, methyl alumoxane, isobutyl aluminumoxane, and tri-n-octyl aluminum, those having bulky or C₆ or higher linear substituents covalently bound to the metal or metalloid center being preferred to minimize adverse interaction with the active catalyst. When an alkyl aluminum or alumoxane is used as activator, any excess over the amount of metallocene present will act as scavenger compounds and additional scavenging compounds may not be necessary. The amount of scavenging agent to be used with metallocene cation-15 noncoordinating anion pairs is minimized during polymerization reactions to that amount effective to enhance activity. In the process described in this invention, it was found that there is an optimum contact time between the scavenger compound and the reaction mixture to maximize catalyst activity. If the contact time is too long, detrimental catalyst deactivation might occur. All documents are incorporated by reference.

The solution process for the production of the EPC elastomers in accordance with this invention improves process economics and increases product capabilities. For process economics, the combination of high catalyst activity and solvent recovery systems brings significant cost improvements. The improved economics of the high catalyst activity solution process compared with the conventional process is related to savings in the finishing area by not requiring catalyst deashing facilities. In addition, efficient solvent recycling also reduces the environmental impact of the process with respect to volatile organic compound emissions to meet increasingly more restrictive regulatory levels. Additionally, the use of the invention process allows for high

diolefin conversion from monomer to incorporated mer unit in the polymer, thus reducing cost of separation and recycle. Typical conversion ratios of diolefin monomer can range from 20%, 30% or up to as high as 40%, and higher.

5 The polymerization process of the invention involves contacting the polymerizable monomers (ethylene, α -olefin and, diene monomer) in solution with the described ionic catalyst composition, preferably at high reaction temperatures, from about 20 °C to 180°C, and can be suitably conducted in the following manner. The solvent is heated to reaction temperature prior to introduction into the reaction vessel.
10 The solvent is then provided to the reaction vessel after polymerizable monomer is introduced in either liquid, gas or solution form into that reaction solvent. A reaction medium is formed comprising the solvent within which the catalyst composition and monomers are contacted for the polymerization reaction. Typically, the scavenging compound is introduced into the reaction solvent to reduce or eliminate catalyst poisons present in any of the reaction medium components prior to introduction into
15 the reactor. If the scavenging compound and activator are different, and contacted with each other for sufficient time, adverse effects on the effectiveness of that activator might occur. In this process, the activator and metallocene compound are contacted in the polymerization reaction vessel in the presence of the polymerizable monomers,
20 comprising the *in-situ* activation.

25 A suitable solution reaction can be conducted at pressures from atmospheric to 500 psig (1-35 bar), preferably from 100 to 300 psig (8 to 21 bar). Preferred reaction temperatures are above 30 °C, preferably up to and above about 80 °C. Typically the polymerization reaction will be exothermic and the reactor or reactor feeds will be chilled or cooled in accordance with known methods to assure that temperatures do not exceed those reasonably suitable for the polymer being produced. Another preferred process in which any of the catalyst, cocatalyst and scavenger selections disclosed in this application can be advantageously practiced is that of a continuous, solution
30 process operated at or above 90 °C to 200 °C, preferably above 110°C. Typically this

process is conducted in an inert hydrocarbon solvent, linear, cyclic or branched aliphatic, or aromatic, at a pressure of from 20 to 200 bar, the reactants being added directly into a suitable reaction vessel containing the solvent at preferred operating temperatures.

5

An additionally suitable homogeneous process of polymerization is that conducted at high pressure, that is at from 200 to 3000 bar, preferably from 500 to 2500 bar in a homogeneous single phase or two fluid phases, with or without unreactive diluents or solvents at temperatures generally above the melting point of the polymer being produced. Such processes are typically known and may include the use of scavenger compounds and catalyst deactivation or killing steps, see for example U.S. patent 5,408,017, WO 95/07941, WO 92/14766, and WO 97/22635. Each of these documents and their U.S. counterparts are incorporated by reference for purposes of U.S. patent practice. Preferred catalyst deactivators, or killers, include high molecular weight, non-recyclable compounds, such as poly vinyl alcohol which exhibit the functional capacity to complex with the catalysts so as to deactivate them while not forming volatile polar by-products or residual unreacted compounds.

The feedstock purification prior to introduction into the reaction solvent follows standard practices in the art, e.g. molecular sieves, alumina beds and oxygen removal catalysts are used for the purification of ethylene, α -olefin, and optional diene. The solvent itself as well, e.g., hexane and toluene, are similarly treated. Purification of the dienes was observed to increase diene conversion, best results were obtained when the diene was fractionally distilled with CaH_2 as the purification method.

25

The α -olefin monomer(s) and diene monomer(s) are introduced in an amount proportional to the levels of incorporation desired for the polymer to be produced and the effective reactive ratios for the polymerizable monomers in the presence of the specific catalyst chosen. In the preferred embodiment the combination of the α -olefin monomer(s) in reaction solvent as introduced into the reactor and effective vapor

pressure of α -olefin monomer(s) is maintained according to the rate of incorporation into the copolymer product. In an alternative embodiment, the partial pressure in the reactor will be provided by ethylene alone in which situation the α -olefin monomer(s) are added solely with reaction solvent. The amounts and vapor pressure will vary according to catalyst selection and polymer to be produced, but can be empirically determined well within the skill in the art, particularly in view of the description provided in the following examples.

The catalyst activator, e.g., non-coordinating anion precursor, ionizing anionic precursor, or alumoxane, can be introduced along with or separately from introduction of the optional diolefin monomer(s), if used. The diolefin can be provided in an amount effective for its rate of reaction and rate of conversion. The catalyst activator can be provided in an amount that is equal to 0.2 to 10 molar equivalents of the Group 4 metallocene compound, preferably 0.25 to 5, and even more preferably 0.33 to 3.0, when a noncoordinating anion precursor. Typically the provision of the noncoordinating anion precursor activator will be in an effective solvent, typically an aromatic solvent such as toluene. Monitoring of polymerization activity by known methods will permit on-line adjustment of alumoxane to assure neither excess nor deficit amounts are maintained for unwanted periods.

-----The scavenging compounds are provided separately afterwards or with one of the foregoing feedstreams, in an amount suitable to increase the activity of the catalyst but in an amount lower than that at which depression of reactivity is observed. Typically an effective amount of the scavenging compound is about 0 (e.g., with an alumoxane activator) to 100 mol. ratio based upon the ratio of scavenging compound to activator, preferably the ratio is 0.3 to 30, and most preferably it is 0.5 to 10.

Ethylene gas is then provided into the reaction vessel in an amount proportional to the level of incorporation desired and the effective reactive ratios for the polymerizable monomers in the presence of the specific catalyst chosen, as with the

5

a-olefin monomer(s). The polymerization starts upon contact of the monomers with the activated catalyst and the rates of supply of each of the components of the composition are adjusted for stable operations at the level of production, molecular weight, monomer incorporation and equipment limitations. The reaction temperature may be permitted to exceed the initial temperature but will preferably be at all times greater than the lower limit of the ranges described above for the invention process.

10

The solvents for the polymerization reaction will comprise those known for solution polymerization, typically the aliphatic solvents represented by hexane, or the aromatic solvents, represented by toluene. Additional examples include heptane, cyclohexane, and Isopar E (C_8 to C_{12} aliphatic solvent, Exxon Chemical Co., U.S.). Preferably the solvent is aliphatic, most preferably it is hexane.

15

Though not strictly necessary for the solution polymerization process as described, the catalyst according to the invention may be supported for use in alternative gas phase, bulk, or slurry polymerization processes where the high activity benefits of the catalysts are sought to be applied. Numerous methods of support are known in the art for copolymerization processes for olefins, particularly for catalysts activated by alumoxanes, any is suitable for the invention process in its broadest scope.

20

See, for example, U.S. patent 5,227,440. An example of supported ionic catalysts appears in WO 94/03056. When using a Lewis acid ionizing catalyst activator a particularly effective method is that described in U.S. patent 5,643,847. A bulk, or slurry, process utilizing supported, *biscyclopentadienyl* Group 4 metallocenes activated with alumoxane co-catalysts is described as suitable for EPM and EPDM in U.S. patents 5,001,205 and 5,229,478, these processes will additionally be suitable with the catalyst compositions of this application. Each of the foregoing documents is incorporated by reference for purposes of U.S. patent practice.

25

30

Though the Examples and the discussion are directed to a single reactor configuration and narrow polydispersity polymers, it is well-known that the use in

series of two such reactors each operated so as to achieve different polymer molecular weight characteristics, or by blending polymers from different reactor conditions or utilizing two or more different transition metal catalysts in one or more reactors, can yield improved processing polymers. The disclosures of U.S. patent 4,722,971 and 5 WO 93/21270 are instructive and are incorporated for purposes of U.S. patent practice. Though directed to the use of vanadium catalysts, the substitution of the catalyst compositions of this invention into one such reactor, or two different invention catalysts into two such reactors, or similar use in two separate polymerizations with subsequent physical blending of the polymer products, will permit tailoring of 10 characteristics (e.g., molecular weights and diene contents) suitable for balancing vulcanization properties with processability. Similarly, the use of mixed catalyst compositions, the invention catalysts with themselves or with others, in one or more such reactors will permit preparation of bimodal or multimodal EPC polymers having improved processing properties.

15

The following examples are presented to illustrate the foregoing discussion. All parts, proportions and percentages are by weight unless otherwise indicated. Although the examples may be directed to certain embodiments of the present invention, they are not to be viewed as limiting the invention in any specific respect.

20 Methods of determining M_n and monomer contents by NMR and GPC for the illustrative EPDM examples of the invention are described in U.S. patent 5,229,478 which is incorporated by reference for purposes of U.S. patent practice. For measurement of comonomer contents in the EPC elastomers, the method of ASTM D3900 for ethylene-propylene copolymers between 35 and 85 wt.% ethylene was used. 25 Outside that range the NMR method was used. See also, U.S. patent 4,786,697 which is incorporated by reference for purposes of U.S. patent practice.

EXAMPLES

Example 1: Synthesis of EPDM

5

The polymerizations were conducted in a 500 cc autoclave reactor operated at the temperature of 115 °C in the batch mode for the polymer and semi-batch for the ethylene monomer. The following procedure was used for the polymerizations:

10

- The reactor was charged with 250 cc of purified hexane, 5 cc of 10 wt.% toluene solution of MAO (activator) and 3 cc of purified ENB (fractional distillation with CaH₂).

15

- The reactor was heated to 115 °C, resulting in a hexane vapor pressure of approximately 37 psig (2.5 bar).
- Propylene was added to the reactor to reach 110 psig (7.48 bar) pressure (liquid phase molar conc. = 0.856M).

20

- Ethylene was added to the reactor to reach 235 psig (16 bar) pressure (liquid phase conc. = 0.871M). These conditions determined the initial ethylene/propylene molar ratio equal to 1.018. The ethylene/ENB molar ratio was 10.43.

25

- The catalyst solution was pumped to the reactor to maintain the polymerization rate constant as indicated by the make-up flow rate of ethylene to the reactor. The pumping was adjusted to keep this rate at about 0.1 SLPM (standard L/min, standard conditions 1 bar, 21.1 °C), to target approximately 10g yield of polymer.

30

- Irganox® 1076 was added to the hexane solution to the final concentration of 0.1 mg/cc to prevent sample degradation. The polymers were worked-up from the solution by precipitation with IPA. After filtering and removing free solvents, the polymer samples were dried under vacuum at 90 °C for about 1 hour.

5

The analysis of the polymers were done by $^1\text{H-NMR}$ for ENB content and GPC for EPDM molecular weight.

Table 1: Results for Example 1

Catalyst: dimethylsilanyl(tetramethylcyclopentadienyl)(cyclopentadienyl)
zirconium dichloride
Activator: methylalumoxane
Temperature : 115 °C
Pressure: 235 psig

10

Polymer Yield (g)	Catalyst Usage (mg)	Ethylene % wt	Propylene % wt	ENB % wt	M_n	PD (MWD)
9.77	0.06	90.77	4.11	3.35	25,974	2.22
13.44	0.02	87.47	4.12	6.79	36,812	1.74

The catalyst efficiency for these two examples were 162.8 and 670 kg-polymer/g-transition metal, respectively.

15

Example 2: Synthesis of EPDM

The same procedure as described in Example 1 was carried out with the noncoordinating anion activator and with the following differences:

20

- At room temperature, after the reactor was charged with 200 cc of hexane, 50 cc of the 1.5×10^{-3} M of the activator solution in toluene was added. The reactor was then heated up to 115 °C, as in Example 1.

5 • To the pressurized reactor after the addition of ethylene, 10 microliters of 2M TIBAL solution in pentane was added as the scavenger at least one minute before the start of the catalyst pumping.

Table 2: Results for Example 2

Catalyst: isopropylidene(cyclopentadienyl)(fluorenyl) zirconium dimethyl
Activator: dimethylanilinium tetra (pentafluorophenyl) borate
Temperature: 115 °C
Pressure: 235 psig

10

Polymer Yield (g)	Catalyst Usage (mg)	Ethylene % wt	Propylene % wt	ENB % wt	M _n	PD (MWD)
14.6	0.1	77	14.55	8.45	80,498	1.65

The catalyst efficiency in this example was 146 kg-polymer/g-transition metal.

Example 3 (Comparative):

Synthesis of EPDM

The same procedure as described in Example 2 was carried out with the following catalysts:

5

Table 3: Results for Comparative Example 2

Catalyst: (see catalyst code following table)
 Activator: (see catalyst code following table)
 Temperature : 115 °C
 Pressure: 235 psig

Catalyst	Polymer Yield (g)	Catalyst Usage (mg)	Catalyst Efficiency (kg/g)*	Ethylene % wt	Propylene % wt	ENB % wt	M _n	PD
C-1	12.8	0.4	32	51.74	33.27	8.15	79,213	1.98
C-1	7.61	0.6	12.68	55.34	29.24	9.06	93,442	1.73
C-2	13	0.12	108.33	68.42	29.09	2.49	76,741	2.04
C-3	6.5	0.01	650	95.94	2.39	0.54	54,239	2.14
C-4	14.12	1.33	10.62	86.77	10.52	2.71	19,314	10.27
C-4	6.59	0.67	9.84	85.21	11.64	3.15	15,853	12.17
C-5	9.4	0.5	18.8	85.14	11.71	3.15	67,723	2.01
C-5	9.05	0.3	30.17	85.43	11.2	3.37	74,125	1.87
C-6	7.39	1.8	4.11	77.65	9.57	9.66	32,559	1.82
C-6	7.96	1	7.96	76.04	10.73	9.84	34,066	1.72

* kg-polymer product per g-transition metal in catalyst

10 C-1: dimethylsilanyl(tetramethylcyclopentadienyl)(cycododecylamido) titanium dichloride/ methylalumoxane
 C-2: dimethylsilanyl(tetramethylcyclopentadienyl)(admantylamido) titanium dimethyl/ dimethylanilinium tetrakis(pentafluorophenyl) borate
 C-3: (pentamethylcyclopentadienyl)(cyclopentadienyl) zirconium dimethyl/ dimethylanilinium tetrakis(pentafluorophenyl) borate

C-4: dimethylsilanyl(2,4-dimethylcyclopentadienyl)(fluorenyl) zirconium dichloride/
methylalumoxane

C-5: dimethylsilanyl(3-n-propylcyclopentadienyl)(fluorenyl) titanium dichloride/
methylalumoxane

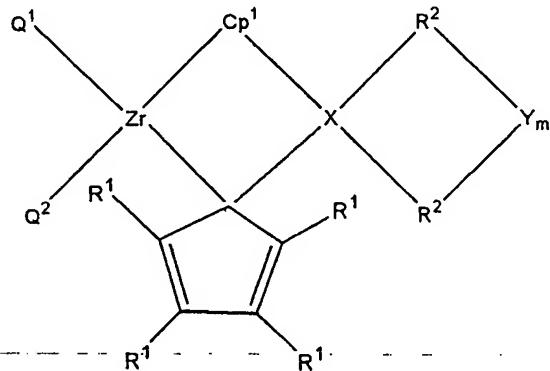
5 C-6: dimethylsilanyl(bis)(indenyl) zirconium dimethyl/ dimethylanilinium tetrakis
(pentafluorophenyl) borate

Comparative Example 3 illustrates that only the polymerization with catalyst C-
3 exhibited the high activity observed for the catalysts according to the invention.
10 However, the polymer prepared with catalyst C-3 exhibited lower incorporation of
comonomers (both propylene and diene monomer) and is accordingly unsuitable in the
process according to the invention.

CLAIMS:

1. A process for the preparation of ethylene copolymers comprising contacting under solution polymerization conditions, ethylene, one or more α -olefin monomer, and optionally one or more diene monomer, with a catalyst composition prepared from a) at least one bridged, bis(cyclopentadienyl) zirconium compound having an unsubstituted cyclopentadienyl ligand, a multiply substituted cyclopentadienyl ligand, said ligands bridged by a covalent bridging group containing one or more Group 14 element, and b) at least one catalyst activator compound.

2. The process of claim 1 wherein said zirconium compound has the following structure:



where the unsubstituted cyclopentadienyl ring is Cp^1 , and the substituted cyclopentadienyl ring has at least two non-hydrogen radical substituents, wherein each R^1 is, independently, a radical selected from:

a) hydrogen radical,

b) hydrocarbyl, silyl or germyl radicals having from 1 to 20 carbon, silicon or germanium atoms,

c) substituted hydrocarbyl, silyl or germyl radicals as defined, wherein one or more hydrogen atoms is replaced by a halogen radical, an amido radical, a phosphido

radical, an alkoxy radical, an aryloxy radical or any other radical containing a Lewis acidic or basic functionality,

d) C₁-C₂₀ hydrocarbyl-substituted metalloid radicals wherein the metalloid is selected from the group consisting of Ge, Sn and Pb,

5 e) halogen radicals,
f) amido radicals,
g) phosphido radicals,
h) alkoxy radicals, and
i) alkylborido radicals; or

10 at least two R¹ groups are joined together to form, along with the carbon atoms to which they are attached, a C₄-C₂₀ ring structure, which is saturated or partially saturated, and substituted or unsubstituted, the ring structure substitution being selected from one or more R¹ groups as defined under (a) – (i) above;

15 X is selected from the group consisting of C, Si, Ge, Sn and Pb;

R² is selected from the group consisting of H, unsubstituted C₁-C₂₀ hydrocarbyl, C₁-C₂₀ hydrocarbyl substituted with at least one Si, and C₁-C₂₀ hydrocarbyl substituted with at least one Ge;

20

m = 0 or 1,

Y is selected from the group consisting of hydrocarbyl radicals, hydrosilyl radicals, and hydrogermyl radicals; and

25

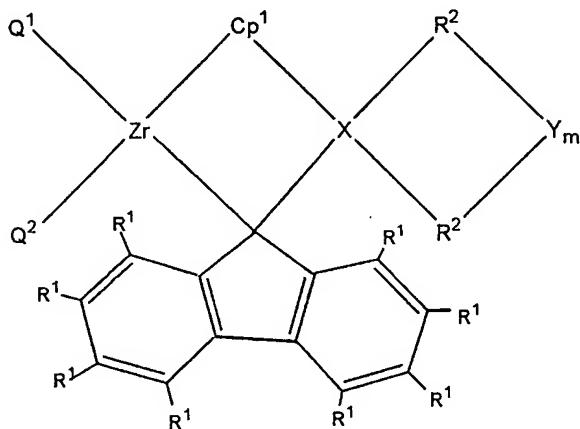
Q¹ and Q² are non-cyclopentadienyl radicals wherein

i) independently, each Q is selected from the group consisting of halide, hydride, unsubstituted C₁-C₂₀ hydrocarbyl, C₁-C₂₀ hydrocarbyl substituted with at least one R¹ group as defined under (a) – (i) above, alkoxide, aryloxide, amide, halide or phosphide; or

30

ii) together, Q¹ and Q² may form an alkylidene, cyclometallated hydrocarbyl or any other divalent anionic chelating ligand.

3. The process of claim 1 wherein said zirconium compound has the following structure:



where the unsubstituted cyclopentadienyl ring is Cp¹, and the substituted fluorenyl ring has at least two non-hydrogen radical substituents, wherein each R¹ is, independently, a radical selected from:

- a) hydrogen radical,
- b) hydrocarbyl, silyl or germyl radicals having from 1 to 20 carbon, silicon or germanium atoms,
- c) substituted hydrocarbyl, silyl or germyl radicals as defined, wherein one or more hydrogen atoms is replaced by a halogen radical, an amido radical, a phosphido radical, an alkoxy radical, an aryloxy radical or any other radical containing a Lewis acidic or basic functionality,
- d) C₁-C₂₀ hydrocarbyl-substituted metalloid radicals wherein the metalloid is selected from the group consisting of Ge, Sn and Pb,
- e) halogen radicals,
- f) amido radicals,
- g) phosphido radicals,

- h) alkoxy radicals, and
- i) alkylborido radicals; or

at least two R¹ groups are joined together to form, along with the carbon atoms to which they are attached, a C₄-C₂₀ ring structure, which is saturated or partially saturated, and substituted or unsubstituted, the ring structure substitution being selected from one or more R¹ groups as defined under (a) – (i) above;

5

X is selected from the group consisting of C, Si, Ge, Sn and Pb;

10

R² is selected from the group consisting of H, C₁-C₂₀ hydrocarbyl, C₁-C₂₀ hydrocarbyl substituted with at least one Si, and C₁-C₂₀ hydrocarbyl substituted with at least one Ge;

m = 0 or 1,

15

Y is selected from the group consisting of hydrocarbyl radicals, hydrosilyl radicals, and hydrogermyl radicals; and

Q¹ and Q² are non-cyclopentadienyl radicals wherein

20

- i) independently, each Q is selected from the group consisting of halide, hydride, unsubstituted C₁-C₂₀ hydrocarbyl, C₁-C₂₀ hydrocarbyl substituted with at least one R¹ group as defined under (a) – (i) above, alkoxide, aryloxide, amide, halide or phosphide; or
- ii) together, Q¹ and Q² may form an alkylidene, cyclometallated hydrocarbyl or any other divalent anionic chelating ligand.

25

4. The process according to Claim 2 wherein the substituted cyclopentadienyl ring is tri-alkyl or tetra-alkyl substituted with methyl, ethyl, isopropyl or tertiary butyl alkyl groups.

5. The process according to Claim 4 wherein the substituted cyclopentadienyl ring is tetramethylcyclopentadiene and A is silacyclobutyl.

6. The process according to Claim 3 wherein the substituted cyclopentadienyl ring is tetramethylcyclopentadiene and A is isopropylidene or diphenylmethylene.

7. The process according to Claim 1-5 wherein said catalyst activator is an alumoxane compound.

10 8. The process according to Claim 1-5 wherein said catalyst activator is an ionizing, anion pre-cursor compound.

15 9. The process of Claim 1-8 wherein said one or more α -olefin monomer is a C₃₋₈ α -olefin and said one or more diolefin monomer comprises 5-ethylidene-2-norbornene or 5-vinyl-2-norbornene.

20 10. The process of Claim 9 wherein said one or more α -olefin monomer is propylene and said one or more diene monomer is 5-ethylidene-2-norbornene or 5-vinyl-2-norbornene.

INTERNATIONAL SEARCH REPORT

Int'l Application No
PCT/US 99/03264

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 C08F210/18 C08F2/06 C08F4/645

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 6 C08F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 729 983 A (MITSUI PETROCHEMICAL IND ;HOECHST AG (DE)) 4 September 1996 see page 4, line 11 - line 15 see page 8, line 17 - line 26 see page 9, line 5 - line 14; claims ----	1-4,6-8
X	WO 97 38019 A (UNIROYAL CHEM CO INC) 16 October 1997 see page 14, line 13 - line 20 see page 15, line 9 - line 10 see page 21, line 29 - page 23, line 18 see page 23, line 19 - line 34; claims; examples ----	1-4,7-10
X	EP 0 612 769 A (TOSOH CORP) 31 August 1994 cited in the application see the whole document ----	1-3,6,8 -/-

Further documents are listed in the continuation-of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

Date of mailing of the international search report

27 May 1999

07/06/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk

Authorized officer

INTERNATIONAL SEARCH REPORT

Inte.inal Application No

PCT/US 99/03264

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X A	EP 0 610 843 A (HOECHST AG) 17 August 1994 see page 7, line 16 - line 20 see page 5, line 54 - page 6, line 5; claims; examples 1-6 ----	1-3,7,9 5
A	EP 0 500 944 A (MITSUI TOATSU CHEMICALS) 2 September 1992 cited in the application see page 5, line 38 - line 49 see page 13, line 29 - line 38 see page 11, line 31 - line 44 see page 6, line 52; claims ----	1-10
A	WO 96 33227 A (EXXON CHEMICAL PATENTS INC) 24 October 1996 cited in the application see page 16, line 7 - line 8; claims ----	1-10

INTERNATIONAL SEARCH REPORT

Information on patent family members

Inte nal Application No

PCT/US 99/03264

Patent document cited in search report	Publication date	Patent family member(s)			Publication date
EP 0729983	A 04-09-1996	JP 8239426 A			17-09-1996
		CA 2170703 A			02-09-1996
		CN 1142511 A			12-02-1997
WO 9738019	A 16-10-1997	EP 0892816 A			27-01-1999
EP 0612769	A 31-08-1994	DE 69406695 D			18-12-1997
		DE 69406695 T			19-03-1998
		JP 7157508 A			20-06-1995
EP 0610843	A 17-08-1994	CN 1096519 A			21-12-1994
		JP 6271627 A			27-09-1994
		US 5733991 A			31-03-1998
EP 0500944	A 02-09-1992	CA 2066247 A			25-01-1992
		DE 69130320 D			12-11-1998
		DE 69130320 T			08-04-1999
		KR 9409020 B			29-09-1994
		EP 0841349 A			13-05-1998
		WO 9201723 A			06-02-1992
WO 9633227	A 24-10-1996	US 5696213 A			09-12-1997
		AU 5557696 A			07-11-1996
		CA 2214454 A			24-10-1996
		CN 1182439 A			20-05-1998
		EP 0821703 A			04-02-1998

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- BLACK BORDERS**
- IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- FADED TEXT OR DRAWING**
- BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- SKEWED/SLANTED IMAGES**
- COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- GRAY SCALE DOCUMENTS**
- LINES OR MARKS ON ORIGINAL DOCUMENT**
- REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.

THIS PAGE BLANK (USPTO)